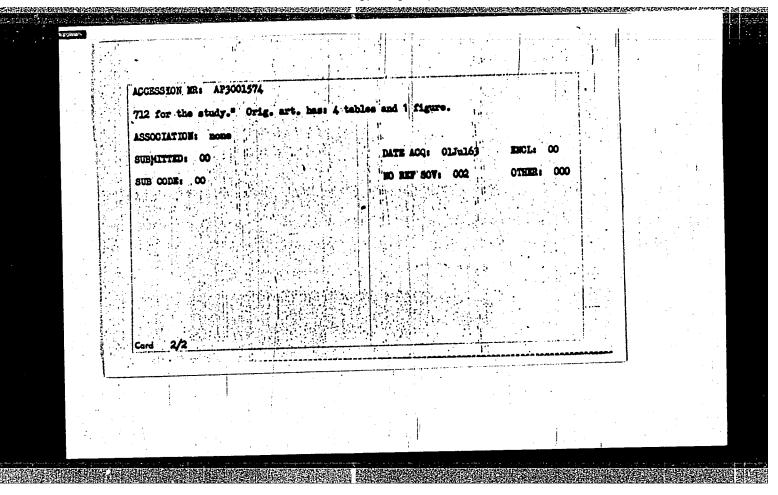


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ACC NR. AP6023429 SOURCE CODE: UR/0190/66/008/007/1219/1225

AUTHOR: Trostyanskaya, Ye. B.; Venkova, Ye. S.; Aristovskaya, L. V.

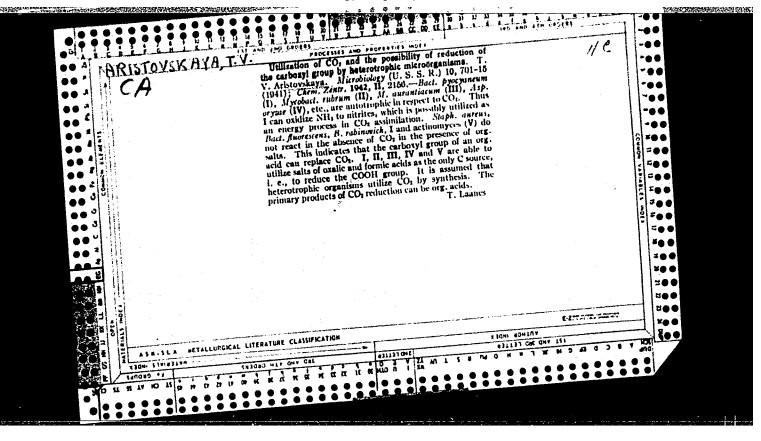
ORG: Moscow Aviation Technology Institute (Moskovskiy aviatsionnyy tekhnologicheskiy institut)

TITLE: Polycondensation of tris(hydroxymethyl)phosphine oxide with phenols

SOURCE: Vysokomolekulyarnyye soyedineniya, v. 8, no. 7, 1966, 1219-1225

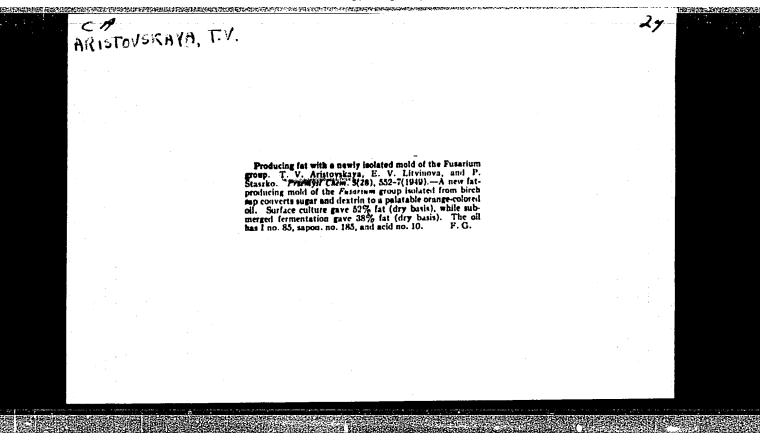
TOPIC TAGS: organic phosphorus compound, alkylphosphine oxide, polymerization rate, polycondensation, catalytic polymerization, heat resistant plastic, thermosetting material, phenolic plastic

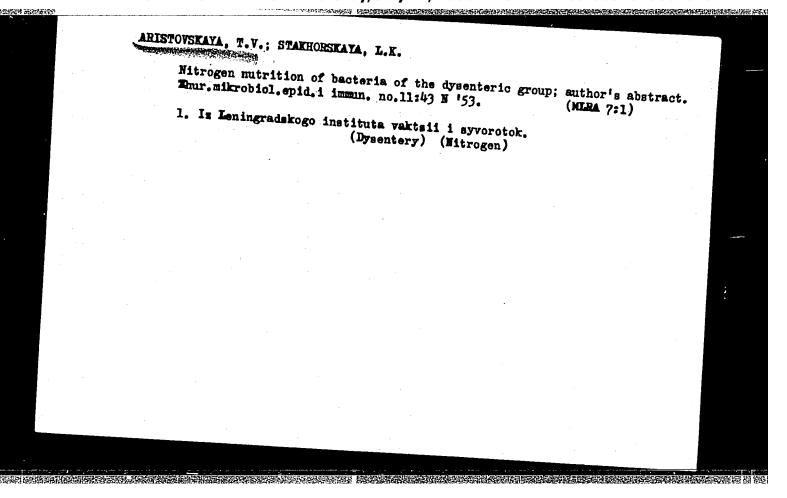
ABSTRACT: Controversial data on the reaction of tris(hydroxymethyl)phosphine oxide (TMPO) with phenol prompted a study of the reactions of TMPO with phenol, resorcinol, and phloroglucinol. The purpose of the study was to prepare phosphorus containing polymers with increased heat and strength resistance. In the absence of a catalyst, the reaction with phenol at 160—200C yielded only a phenol-formaldehyde type resin with a low polycondensation rate because of predominant splitting of TMPO. In the presence of a BF3-phenol complex as catalyst, the rate of polycondensation was increased and at 145C in 8 hr the reaction yielded 30 wt% of an insoluble, thermosetting phenol-TMPO resin with high phosphorus content. The polycondensation rate was further increased in the reaction with resorcinol and even more so with phloroglucinol. Chemical analysis of the reaction products at various stages of polycondensation indicated Card 1/2

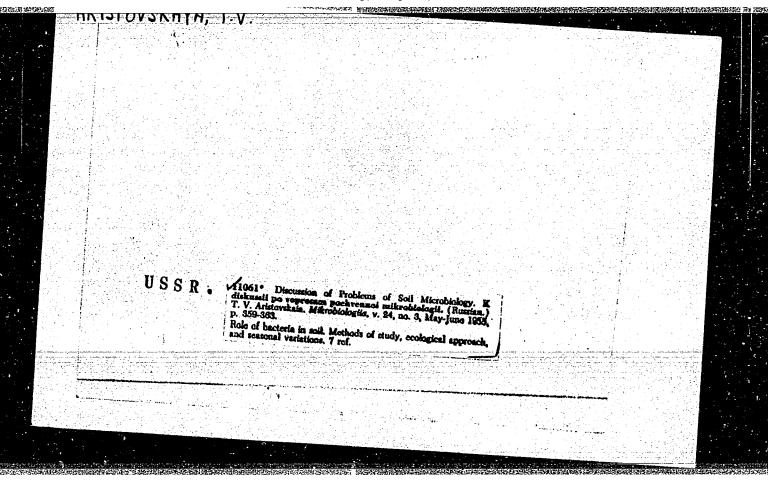


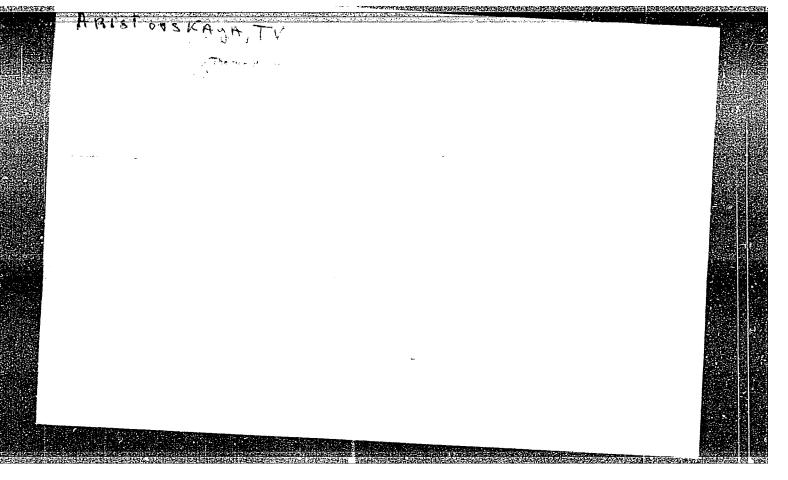
ARISTOVSKAYA, T. V.	
"Carbon Dioxide and Aristovskaya, T. V.	the Life of Heterotrophic Microorganisms (p. 54) by (Moscow).
SO: Advances in Con	(Moscow). ntemporary Biology (Uspekhi Sovremennoi Biologii) Vol. 17, 1944, No. 1
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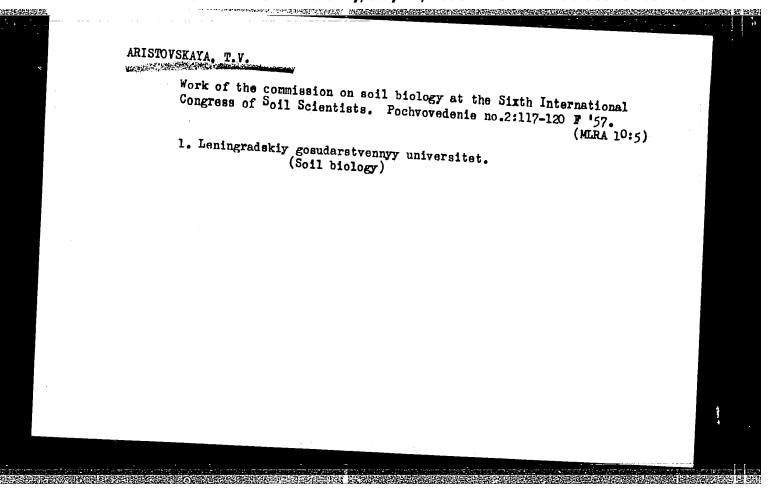
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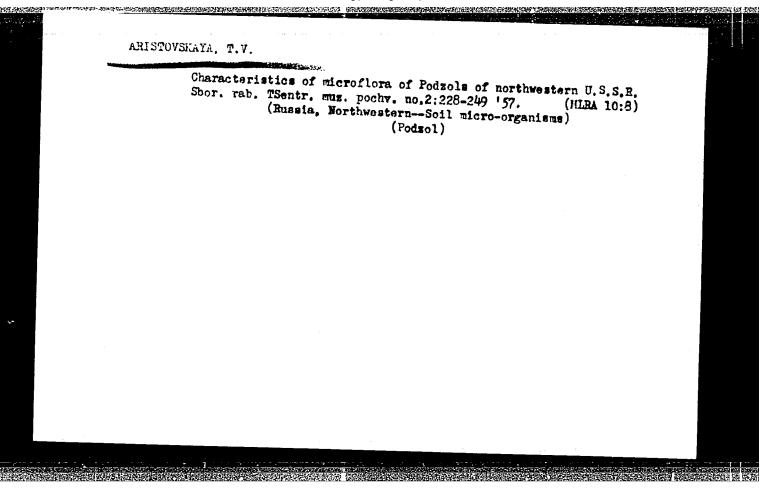








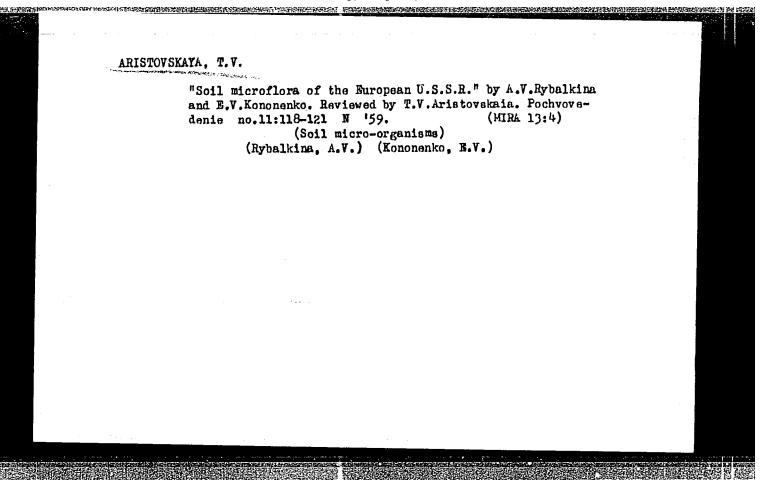


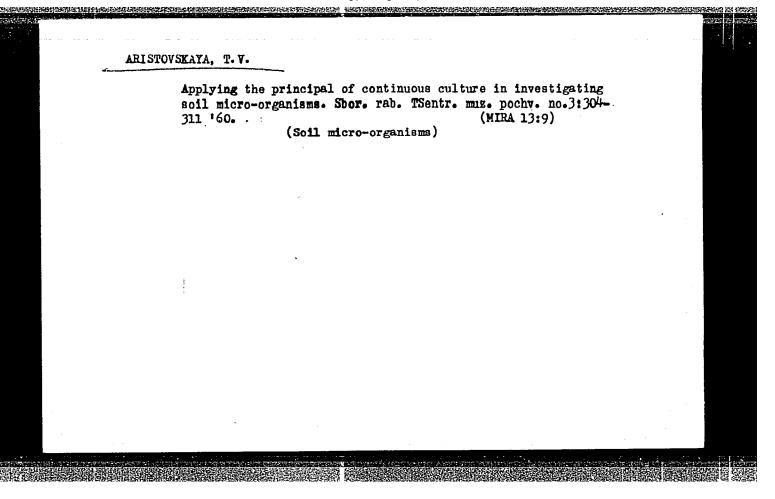


Decomposition of fulvic acid by micro-organisms [with summary in English]. Pochvovedenie no.11:40-51 N '58. (MIRA 11:12) 1. TSentral'nyy Muzey Pochvovedeniya imeni V.V.Dokuchayeva AN SSSR. (Fulvic acid) (Soil micro-organisms)

Seasonal variations and ecological characteristics of microflora in some Podzol soils. [with summary in English]. Mikrobiologida 27 no.3:324-330 My-Je '58 (MIRA 11:9)

1. TSentral'nyy muzey pochvovedeniya im. V.V. Dokuchayeva AN SSSR, Leningrad. (SOIL, microbiology seasonal & ecol. factors (Rus))

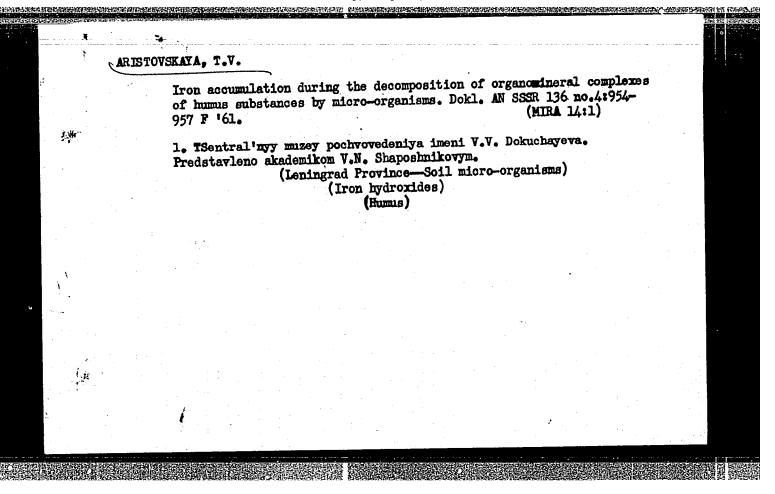




ARISTOVSKAYA, T.V.; PARINKINA, O.M.

New methods of investigating communities of soil micro-organisms.
Pochvovedenie no.1:20-28 Ja '61. (MIRA 14:1)

1. TSentral'nyy muzey pochvovedeniya imeni V.V. Dokuchayeva AN SSSR. (Soil micro-organisms)



ARISTOVSKAYA, T.V.; PARINKINA, O.M.

Studying soil microbe patterns of Leningrad Province. Mikrobiologiia 31 no.3:385-390 My-Je '62. (MIRA 15:12)

1. TSentral'nyy muzey pochvovedeniya imeni V.V.Dokuchayeva, Leningrad.

(LENINGRAD PROVINCE -- SOIL MICRO-ORGANISMS)

ARISTOVSKAYA, T.V.; VI.ADIMIRSKAYA, M.Ye.; COLLERBAKH, M.M.; KATANSKAYA,

F.A.; KASIKIN, P.N.; KLUPP, S.Ye.; LOZINA-LOZINSKIY, L.K.; NORKINA,

S.P.; RUMYANTSEVA, V.M.; SELIBER, G.L., prof. [deceased]; SKALON, {
 I.S.; SKORODUMVA, A.M.; KHETAGUROVA, F.V.; CHASTUKHIN, V.Ya.;
 PARSADANOVA, K.G., red.; GARINA, T.D., tekhn. red.

[Comprehensive laboratory manual on microbiology] Bol'shoi praktikum po mikrobiologii. [By] T.V.Aristovskaia i dr. Pod obshchei red. G.L.Selibera. Moskva, Vysshaia shkola, 1962. 490 p.

(MICROBIOLOGY—LABORATORY MANUALS)

(MICROBIOLOGY—LABORATORY MANUALS)

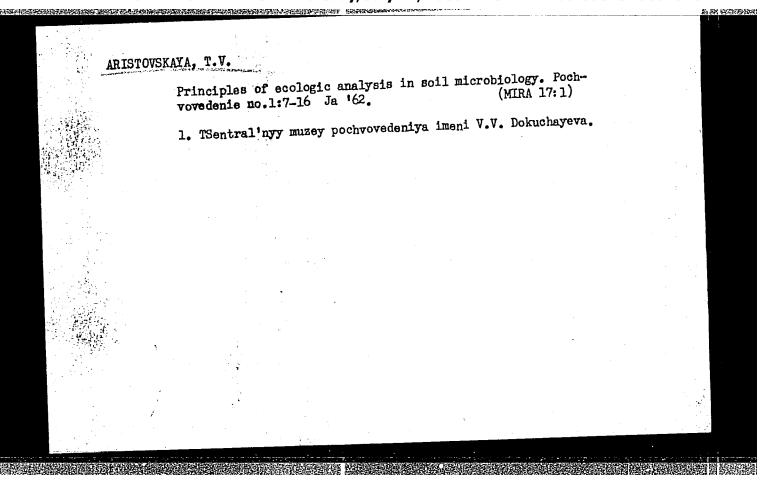
Decomposition of organic-mineral compounds in Podzolic soils. Pochvovedenie no.1:30-43 Ja '63. (MIRA 16:2) 1. TSentral'nyy muzey pochvovedeniya imeni V.V.Dokuchayeva. (Podzol) (Humus) (Soil micro-organisms)

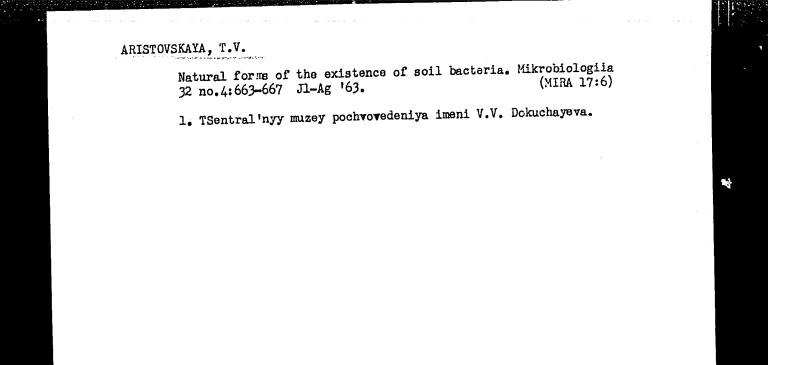
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New soil microorganism Seliberia stellata noc.gen.n.sp.
Izv. AN SSSR. Ser. biol. 28 no. 1:49-56 Ja-F'63. (MIRA 16:8)

1. TSentral'nyy muzey pochvovedeniya imeni V.V.Dokuchayeva,
Leningrad.

(SCIL MICROORGANISMS)

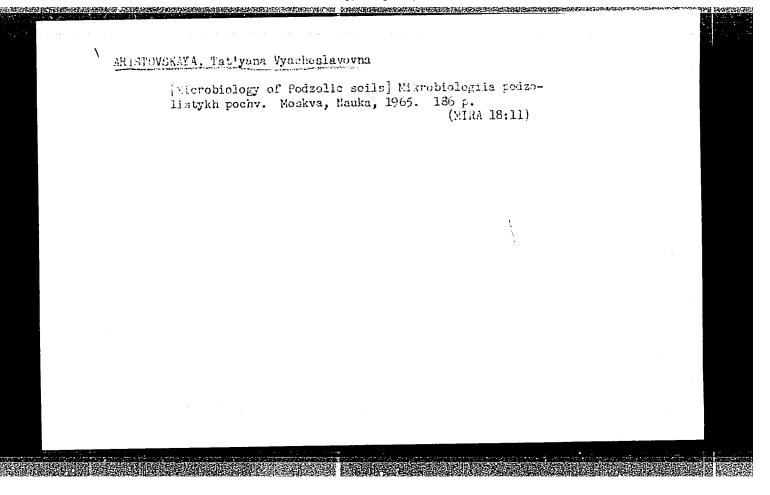


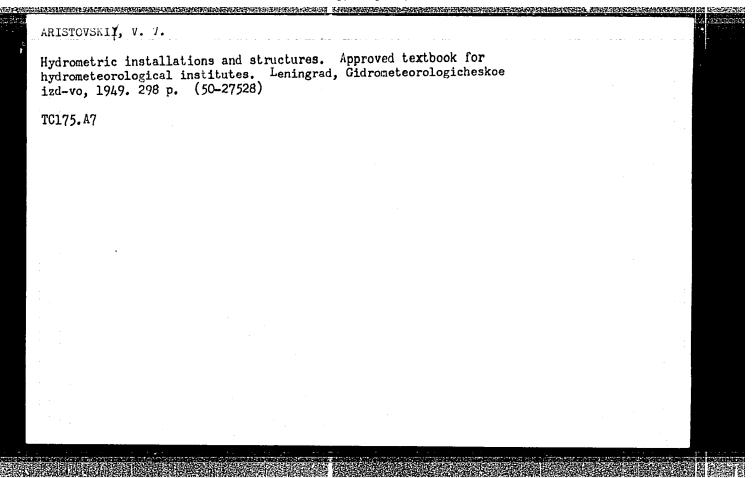


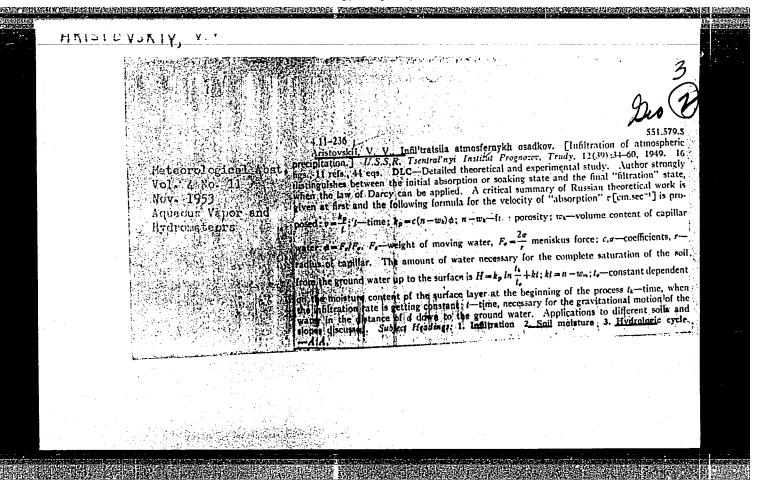
Taxonomic position of the genus Seliberia Arist. et faring.
Mikrobiologiia 33 no.6:929-934 N-D '64.

1. TSentral'nyy muzey pochvovedeniya imeni Dokuchayeva.

(NIRA 18:4)







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PISHKIN, B.A. [Pyshkin, B.A.], otv.red.; ARISTOVSKIY, V.V. [Aristovs'kyi, V.V.], doktor tekhn.nauk, red.; GUZOV, M.Z. [Huzov, M.Z.], kand.tekhn.nauk, red.; ZAGUMKNNYY, O.G. [Zahumennyi, O.H.], red.; PECHKOVSKAYA, O.M. [Piechkovs'ka, O.M.], red.izd-va; MIL'OKHIN, I.D., tekhn.red.

[Calculation of seepage through hydraulic structures; collection of scientific works] Fil'tratsiini rozrakhunky gidrotekhnichnykh sporud; zbirnyk naukovykh prats'. Kyiv, 1959. 161 p.

1. Akademiia nauk URSR, Kiev. Rada po vyvchenniu produktyvnykh syl URSR. 2. Chlen-korespondent AN URSR, golova Komisii po problemi kompleksnogo vikoristannya vodnikh resursiv URSR RPS AN URSR (for Pishkin).

(Hydraulic engineering--Tables, calculations, etc.)

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PISHKIN, B.A. [Pyshkin, B.A.], otv.red.; TYULENEV, M.O. [Tiuleniev, M.O.], red.; ARISTOVSKIY, V.V. [Aristovs'kyi, V.V.], doktor tekhn.nauk, red.; ALPAT'YEV, S.M. [Alpat'iev, S.M.], kand. sel'skokhoz.nauk, red.; ZHELEZNYAK, Y.A. [Zheliezniak, I.A.], kand.tekhn.nauk, red.; MAKSIMCHUK, V.L. [Maksymchuk, V.L.], kand.tekhn.nauk, red.; SEMENOV, K.S., kand.tekhn.nauk, red.; PECHKOVSKAYA, O.M. [Piechkovs'ka, O.M.], red.izd-va; KADASHEVICH, O.O., tekhn.red.

[Over-all utilization of Ukrainian water resources; collected studies] Komplekane vykorystannia vodnykh resursiv Ukrainy; sbirnyk naukovykh prats'. Kyiv, 1959. 173 p. (MIRA 13:1)

1. Akademiia nauk URSR, Kiev. Rada po vyvchenniu produktyvnykh syl URSR. 2. Chlen-korespondent AN URSR; golova Komisii po problemi kompleksnogo vikoristannya vodnikh resursiv URSR, Rada po vivchennyu produktivnikh sil URSR Akademii nauk URSR (for Pishkin).

3. Chlen-korespondent AN URSR; Ukrains'kiy naukovo-doslidniy institut gidrotekhniki ta melioratsii (for Tyulenev). 4. Institut gidrologii i gidrotekhniki AN URSR (for Zheleznyak, Maksimchuk, Pishkin).

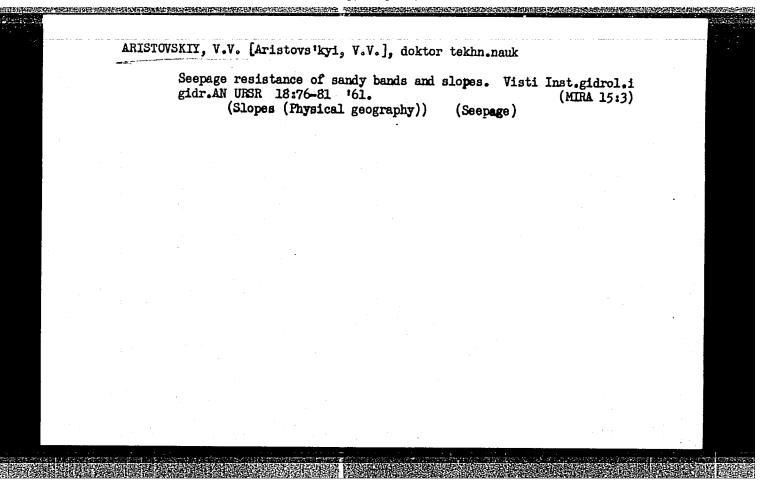
(Ukraine--Water resources development)

PYSHKIN, B.A., red.; ARISTOVSKIY, V.V. [Aristovs'kyi, V.V.], doktor tekhn.
nauk, red.; DYATLOVITSKIY, L.I. [Diatlovyts'kyi, L.I.], kand. tekhn.
nauk, red.; SPIRIN, G.M. [Spirin, H.M.], red.; SPIRINA, N.I., red.;
PECHKOVSKAYA, O.M. [Pechkovs'ka, O.M.], red. izd-va; RAKHLINA, N.P.,
tekhn. red.

[Investigating the stressed state of hydraulic structures] Doslidzhennia napruzhennoho stanu hidrotekhnichnykh sporud; sbirnyk naukovykh prats'. Kyiv, 1961. 149 p. (MIRA 14:10)

1. Akademiya nauk URSR, Kiev. Rada po vyvchenniu produktyvnykh syl URSR. 2. Chlen-korrespondent AN URSR (for Pyshkin).

(Hydraulic structures)



PYSHKIN, B.A., otv. red.; ARISTOVSKIY, V.V., doktor tekhn.nauk, prof., red.; RUSAKOV, S.V., kand. tekhn. nauk, red.; MAKSIMCHUK, V.L., kand. tekhn. nauk, red.; TSAYTS, Ye.S., kand. tekhn. nauk, red.; PECHKOVSKAYA, O.M., red.; LIBETMAN, T.R., tekhn. red.

[Changes in the banks of reservoirs]Pererabotka beregov vodokhranilishch. Kiev, Izd-vo Akad. nauk USSR, 1962. 140 p. (MIRA 15:11) 1. Akademiya nauk URSR, Kiev. Rada po vyvchenniu produktyvnykh

1. Akademiya nauk URSR, Kiev. Rada po vyvchenniu produktyvnykh syl. 2. Chlen-korrespondent Akademii nauk Ukr. SSR (for Pyshkin). (Reservoirs) (Coast changes)

ARISTOVSKIY, Valer'yan Valer'yanovich [Arystovs'kyi, V.V.], doktor tekhm. nauk; SLOBODYAN, Roman Tikhonovich, kand. tekhm. nauk; DIDKOVSKIY, M.M.[Didkovs'kyi, M.M.], kand. tekhm. nauk, otv. red.; REVERA, O.Z., kand. geogr. nauk, nauchnyy red.; DAKHNO, Yu.M., tekhm. red.

[Stability of the Kakhovka Reservoir shores undergoing deformations caused by subsidences and slides]Stiikkist' berehiv Kakhovs'koho vodoskhovyshcha, shcho zaznaiut' szuvnykh ta prosadochnykh deformatsii. Kyiv, Vyd-vo Akad. nauk URSR, 1962. 145 p. (MIRA 15:11)

(Kakhovka Reservoir--Coast changes)

ARISTOVSKIY, V.V., doktor tekhn.nauk, prof.; TSERAPNIYER, L.S., ingh.;

LAPINE, L.V., ingh.; YEFREMOVA, Ye.A., ingh.

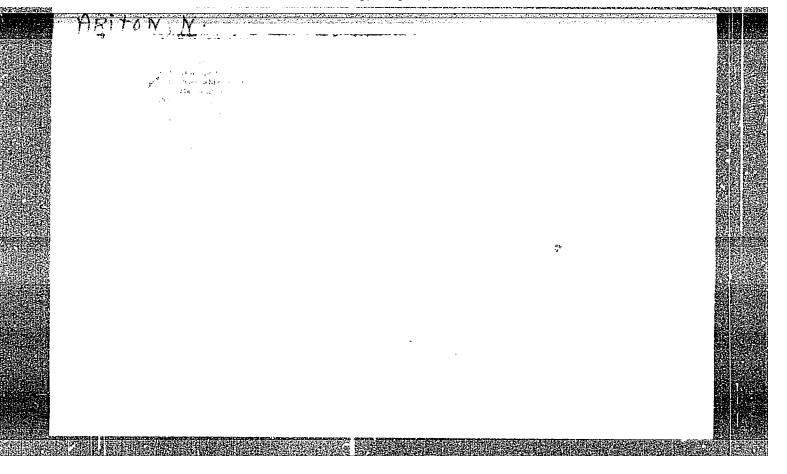
"German-Russian hydraulic engineering dictionary" edited by M.M.

Grishin. Reviewed by V.V.Aristovskir and others. Gidr. stroi.
33 no.5:62-63 My '63.

(Hydraulic engineering—Dictionaries)

(German language—Dictionaries—Russian) (Grishin, M.M.)

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ARITON, N.

RUMANIA/Analytical Chemistry - Analysis of Inorganic Substances. E.

E-2

Abs Jour

: Ref Zhur - Khimiya, No 8, 1958, 24775

Author

: Popper, E., Ariton, N., Popa-Craciuneanu, R.

/ Inst

: Rumanian Academy.

Title

Rapid Semi-Micro Method of Gravimetric Determination of

Lead.

Orig Pub

Studii si cercetari chim. Acad. RPR Fil. Cluj, 1956, 7,

No 1-4, 85-88

Abstract

In the synthesis of 2-mercapto-5-anilono-1,3,4-thiodiazole used in the determination of Pb²⁺ and Hg²⁺(RZhKhim, 1958, 24757, 24774), there is obtained as an intermediate product the anilide of hydrazine-N,N'-bis-thiocarboxylic acid (I), which is a white crystalline substance, MP 192-193°, readily soluble in pyridine, acetone and alkalies,

less soluble in C2H5OH and insoluble in water.

Card 1/2

7

RUMANIA/Analytical Chemistry - Analysis of Inorganic Substances.

E-2

Abs Jour : Ref Zhur - Khimiya, No 8, 1958, 24775

I precipitates quantitatively Pb in the form of a yellow crystalline precipitate which is suitable for a gravimetric determination of Pb. To 0.5-5 ml of the solution being analyzed, containing not more than 200 //ml Pb; are added about 20 ml C₂H₂OH, 0.2% alcoholic solution of I sic and the mixture is allowed to stand for 1 hour. The resulting precipitate is filtered off, washed, first with alcohol then with ether, dried in a vanuum desiccator, and weighed. I precipitates also Ag + Hg + +, Hg + +, Bi + +, Cu + +, Co + +, Mn + + and Cr + +. Alkali and alkaline earth metals do not interfere. Duration of determination of Pb is 1.5-2 hours; error ± 0.4%.

Card 2/2

ARITON, N.

RUMANIA/Analytical Chemistry - Analysis of Chorganic Substances. E-2

: Ref Zhur - Khimiya, No 8, 1958, 24757 Abs Jour

Pepper, E., Ariton, N., Proinov, L., Craciuneanu, R. Author

Inst

New Rapid Method of Gravimetric Determination of Mercury Title

: Rev. chim., 1957, 8, No 9, 594-596 Orig Pub

: For the determination of Hg2tuse is made of a new reagent Abstract

2-mercapto-5-anilino-1,3.4-thiodiazole (I) (RZhKhim, 1958, 24774), which is a white-yellow powder, MP 215-216°, readily soluble in C2H5OH, less soluble in alkalies and insoluble in water. I precipitates Ag⁺, Pb²⁺, Hg³⁺, Hg³⁺, Bi³⁺, Cu³⁺, Co³⁺ and Fe³⁺. Hg²⁺ ions form with I a yellow precipitate insoluble in C_M_OH and in ether and suitable for gravimetric determination of Hg. The solution being analyzed, containing Hg as Hg Cl , or Hg (NO ,) (concentra-

tion of free HNO3 not above 0.2-0.3 N), is diluted with twice its volume of C₂H₃OH, and Hg²⁺is precipitated with

Card 1/2

RUMANIA/ Analytical Chemistry - Analysis of Inorganic Substances. E-2
Abs Jour : Ref Zhur O Khimiya, No 8, 1958, 24757

a 5% alcohol solution of <u>I</u> under continuous stirring. Solution and precipitate are heated on a water bath until the amorphous precipitate becomes crystalline, filtered immediately through a filter crucible No 5 or No 4, the precipitate is washed with alcohol and ether, dried in a vacuum-desiccator and weighed. If the concentration of HNO in the solution being analyzed is above 0.3 N the solution is evaporated to dryness and the residue is dissolved in water acidified with HCl. The minimum determinable amount of Hg³⁺ is 40/ml. Duration of determination is from 45 minutes to 2 hours; error ± 0.54%. The method is suitable for determination of Hg in medicinals.

Card 2/2

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	COUNTRY		
	ABS. JOUR.	: RZKhim., No. 21 1959, No. 75793	
	ROHTUA	Popper, E., Ariton, N., Proinov, L., and Craciunea-	
	INST. TITLE	Not given The Analysis of Mercury Medicinal Preparations by	
	ORIG. PUB.	Anilino-1,3,4-Thiodiazore Farmacia (RPR), 6, No 6, 491-498 (1958)	
	ABSTRACT	A new gravimetric is proposed for the determina- tion of Hg in medicinal preparations, using 2-	
	Maria Nakaja ka	mercapto-5-anilino-1.5, to the calculate indicated reagent is very sensitive (accurate indicated reagent is very sensitive (accurate determinations of Hg in medicines are possible determinations of Hg of 40 gammas per	
	, kurt dár – i	ml), it is readily prepared. The authors propose any analytical laboratory. The authors propose	
		for the official methods in the Rumanian Pharmaco- ber of medicines listed in the Rumanian Pharmaco-	
	CARD: 1/1	poeia. From authors' summary nu, R.	·
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Li i			

"APPROVED FOR RELEASE: Thursday, July 27, 2000

CIA-RDP86-00513R00010201

ACC NR: AP7005131

(A)

SOURCE CODE: UR/0126/66/022/004/0556/0562

AUTHOR: Yegiyan, K. A.; Arityunyan, R. G.

ORG: none

TITLE: Coercive force of thin ferromagnetic films as a function of certain factors

SOURCE: Fizika metallov i metallovedeniye, v. 22, no. 4, 1966, 556-562

TOPIC TAGS: magnetic thin film, ferromagnetic film, iron nickel alloy, magnetic coercive force

ABSTRACT: The nature of the coercive force $H_{\rm c}$ of thin films is as yet inadequately understood, and previous studies of $H_{\rm c}$ investigated its dependence on only some one factor or another rather than on several factors. To fill this gap, the authors investigated the dependence of $H_{\rm c}$ on a number of factors: composition of binary Fe-Ni alloy film (varying from 75% Ni and 25% Fe to 84% Ni and 16% Fe); dependence of $H_{\rm c}$ on substrate temperature (150-350°C); dependence of $H_{\rm c}$ on film thickness (700-2700 Å); dependence of $H_{\rm c}$ on the anisotropy field $H_{\rm c}$ and its angular dispersion ϕ . Only uniaxially anisotropic films were considered. The films were obtained by the vacuum evaporation method in 1-2·10⁻⁵ mm Hg at the rate of 120-160

Card 1/2

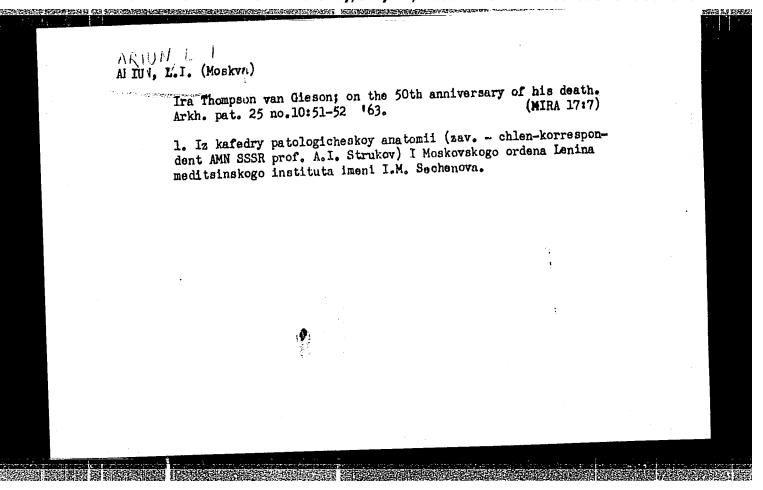
UDC: 539.216.2:538.248

ACC NR: AP7005131

A/min, with deposition on ordinary glassslides. Hysteresis properties were measured with the aid of a ferrotester at a frequency of 1200 cps. H_c was determined in a field of 5 oe; H_k, by the extrapolation method; and ϕ , by the Krouter pulsed method. The magnetoelastic constant was determined according to the dependence of H_k on relative elongation. Findings: under specific experimental conditions there exists a direct relationship between the variations in H_c and angular dispersion. E.g. the minima of both these quantities coincide for the alloys with the composition 75% Ni +25% Fe. For films 700 to 2700 Å thick $H_c \sim d^{-2/3}$. The increase in ϕ with increase $\,$ in film thickness in this case is associated with the increase in the demagnetizing field of the specimen. The mechanism of action of such angular dispersion on the properties of the films differs from that of the conventional structural increase in φ . In particular, in this case a rise in φ leads to a decrease rather than increase in H_c . The anisotropy field H_k contributes to the magnitude of H_c only in the region $H_c/H_k > 0.8$. When $H_c/H_k < 0.5$ the variation in H_k does not lead to variation in H_c . It is assumed that this relationship is conditioned by the change in the mechanism of the reversal of films along the easy axis with increase in the ratio H_C/H_k. "The authors are indebted to Professor R. V. Telesnin for discussion of this project." Orig. art. has: 7 figures.

SUB CODE: 420/ SUBM DATE: 08Jan66/ ORIG REF: 003/ OTH REF: 009

Card 2/2



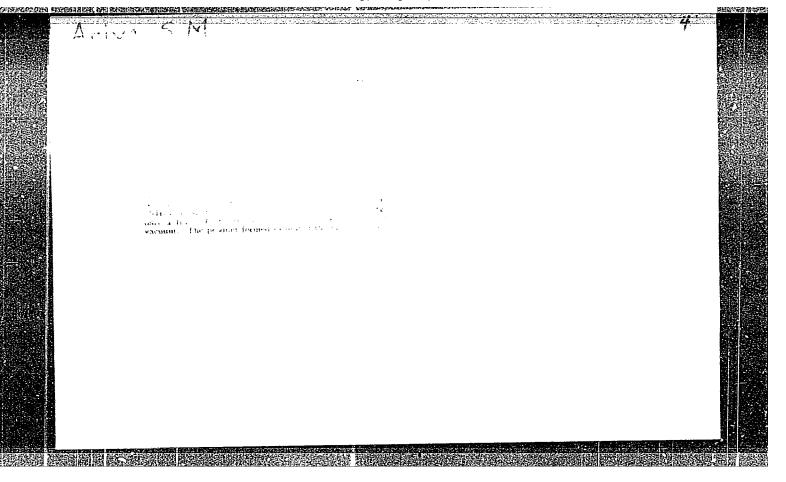
ARIYA, S. M.

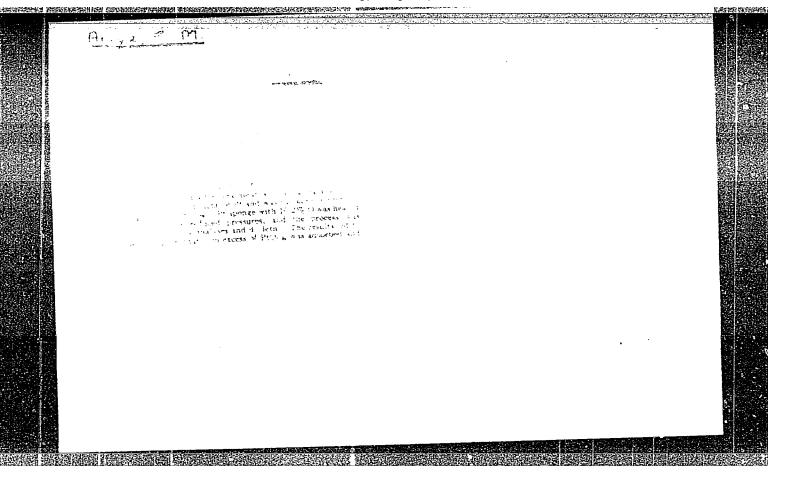
Annual Scientific Session of Leningrad University in 1951, P.G.Makarov, T.A. Agekyan, G. Drakarev, N. Yanovskaya, S.V.Golodnikov, and S.M.Araya, Vest Leningrad U, Ser Mat, Fiz, Khim, Vol 7, no 2, pp 184-196, Feb 52.

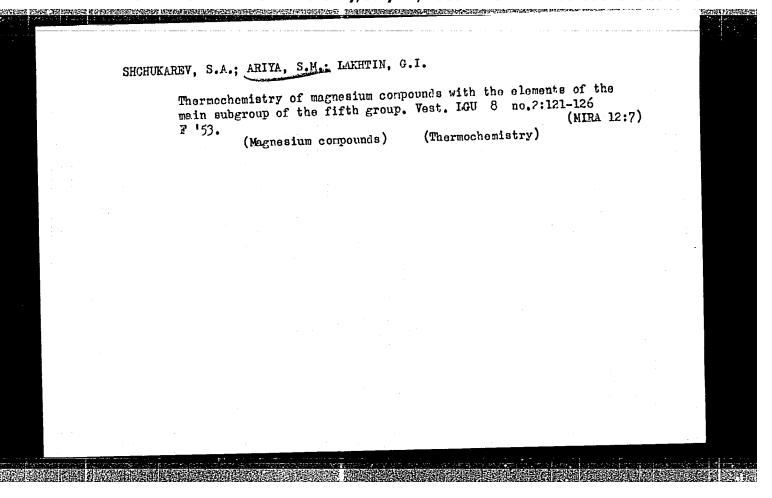
The annual scientific session of Leningrad University took place 4-20 February 1952. The Math Section was subdivided into math, mechanics, and astronomy; the physics comprosed also geophysics. The Chemistry Section dealt also with cooperation with industry.

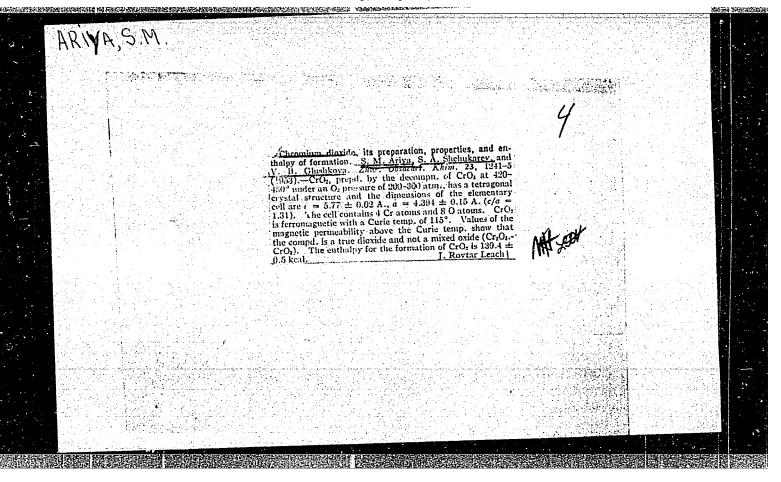
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i ee	kii zakon prirody (Great law of nature). Periodicheskaia sistema elementov Mendeleeva znachenie v nashi dni. Moskva, Detgiz, 1953. 111p.													
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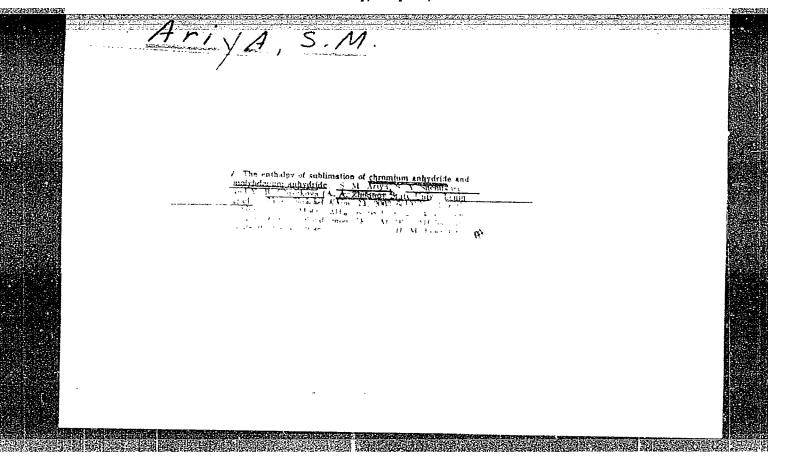








ARIYA, S. M.	χ H ,	Khim, Vol 23, No 9, pp 1455-1450 a calorimetric detn of the enthalpy of Aptor 307 + 1.39(H2) = \begin{align*} \begin{align*} \pi \eqrig \pi \eqrig \eqrig \pi \eqrig \eqr	2borr25 the of	268T25	
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	is Latinu Le Ple Doca al eningi	, pp n of Hg) = invol ound:	igati diss		
	USSR/Chemistry - Platinum Oxides "Investigation of the System Platinum-Oxygen. Enthalpy of the Formation of the Platinum Oxide Pt304, "S.M. Ariya, M.P. Morozoca and A.A. Reykhardt, Chair of Inorg Chem, Leningrad State U.	Zhur Obshch Khim, Vol 23, No 9, pp 1455-1450 By means of a calorimetric detn of the enthalpy of the reaction \sqrt{P} to, $3\sqrt{7} + 1.39(R_2) = \sqrt{P}t/7 + 1.39R_2$ the magnitude of the enthalpy involved in the forther mation of platinum oxide was found: $\Delta H_F PtO_{1.3} G_{-13.6}$ 1.1 kcal. The indicated magnitude agrees	results of an investigation ium of the process of dissocovide.		
	Syst Syst M.P. N	Zhur Obshch Khim, Vol 23, No 9 by means of a calorimetric det the reaction \(\begin{align*}{c} \frac{\text{T}}{30} \end{align*} + 1.39 \\ \text{the magnitude of the enthalpy mation of platinum oxide was 1-13.6 \(\frac{\text{T}}{1}\). It keal. The indicates	an inve		
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ARIYA, S.M.

USSR/Chemistry - Physical chemistry

Card 1/1

Pub. 151 - 7/36

Authors

Ariya, S. M.; Morosova, M. P.; and Shneyder, L. A.

Title

Thermodynamics of oxide phases of various composition. Part 1.- On the thermodynamics of FeO

Periodical

Zhur. ob. khim. 24/1, 41-47, Jan 1954

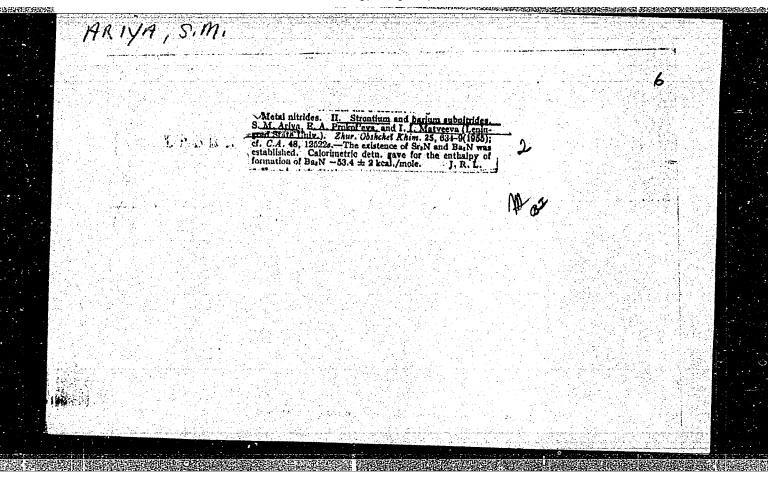
Abstract

The equilibrium of the FeO reduction process with CO2/CO mixtures was investigated at 1104 and 1182°K. The constant of the equilibrium process in the investigated temperature range was found to be practically independent from the temperature in zones of FeO homogeneity. It was established that the dependence of the equilibrium pressure of atomic oxygen upon the composition of the solid phase does not respond to the Henry law and that the entropy of FeO, computes per 1 g/atom, increases somewhat in accordance with the increase in oxygen content of that particular phase. The enthalpy values for the formation of various types FeO, at standard conditions, were determined. Six references: 3-USA; 2-German and 1-USSR (1922-1949). Tables; graphs.

Institution: The A. A. Zhdanov State University, Leningrad

Submitted

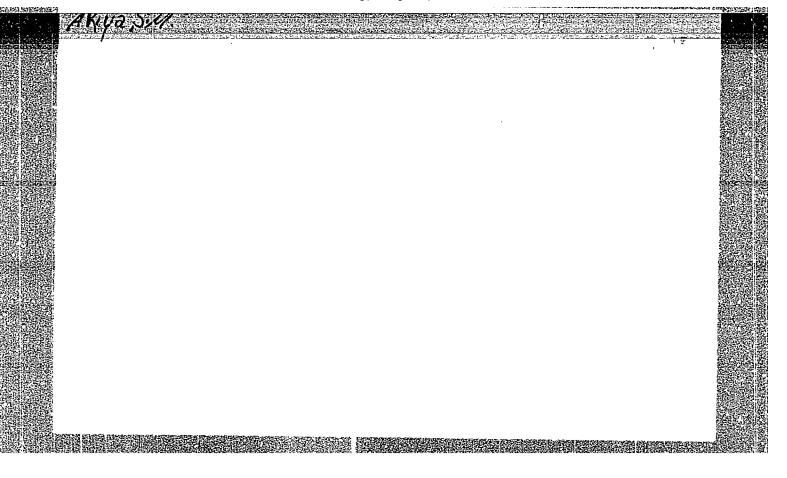
August 8, 1953

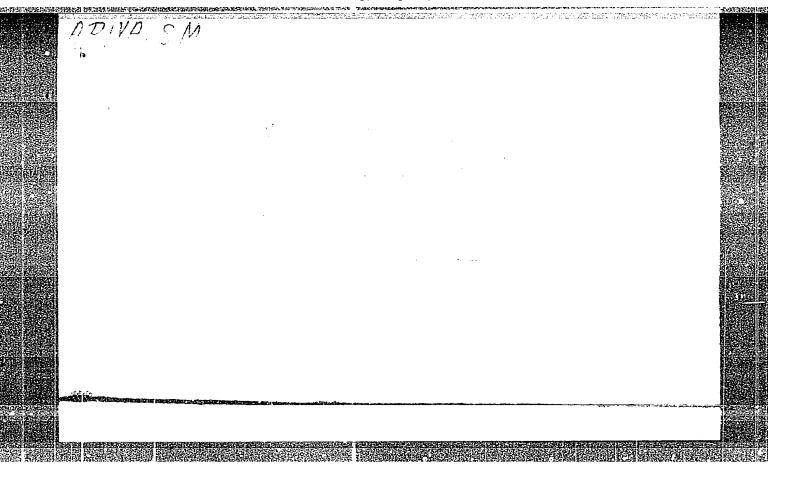


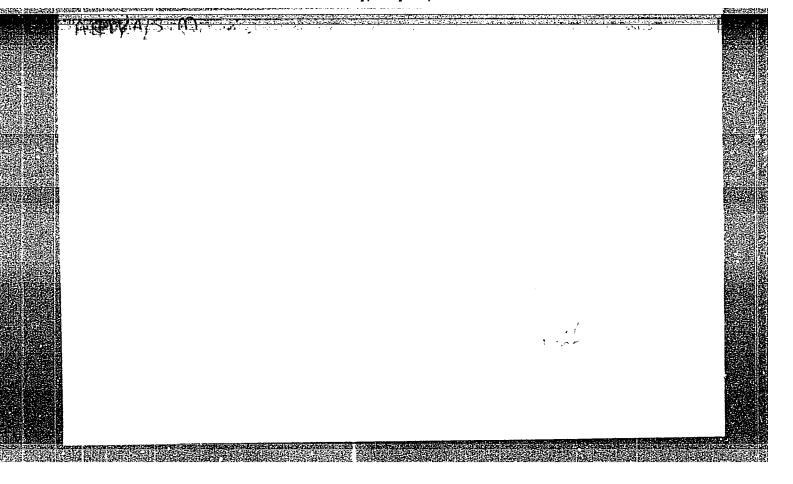
ARIYA,S.M.; PROKOF'YEVA,Ye.A.

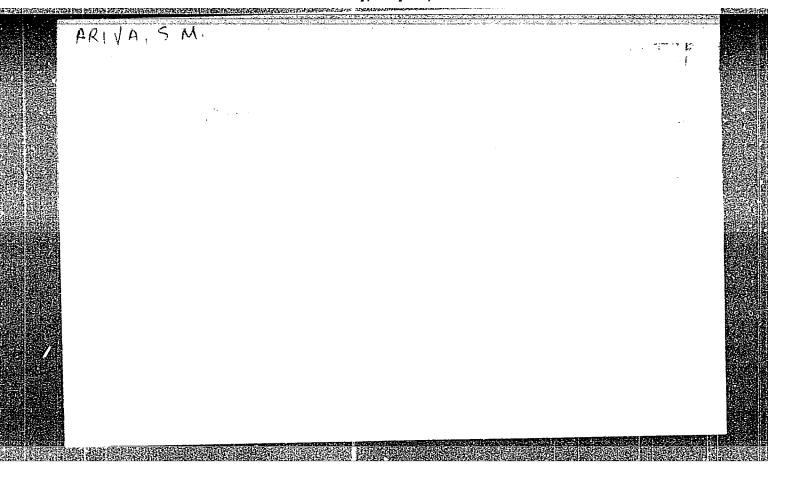
Investigation of metal nitrides. Part 3. Phase diagram of Ba-N
systems in the high-pressure range. Zhur.ob.khim.25 no.5:849-851
My'55. (MIRA 8:10)

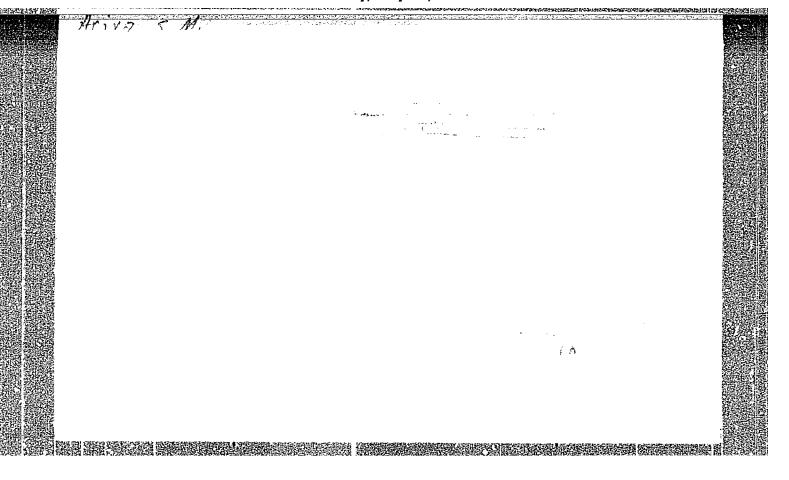
1. Leningradskiy Gosudarstvennyy universitet
(Barium compounds) (Nitrides)











ARIYA, S.M.; ZASLAVSKIY, A.I.; MATVEYEVA, I.I.

Chemistry of the compounds of a variable composition. Part 4.

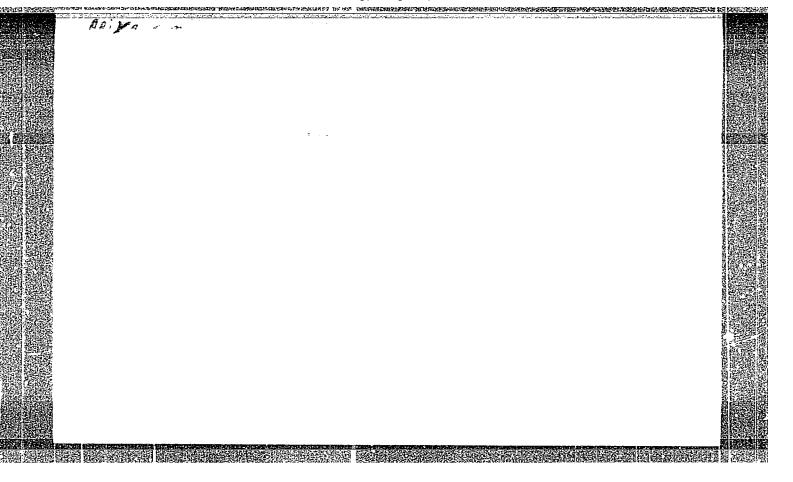
System tantalum -- selenium. Zhur.ob.khim. 26 no.9:2373-2375

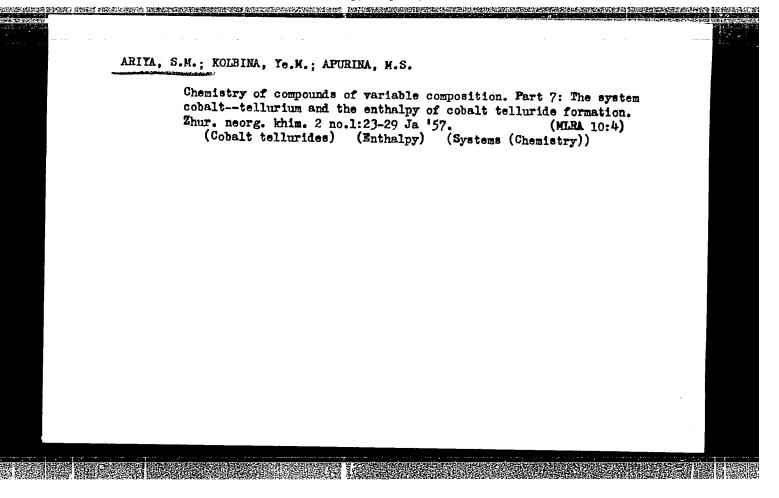
S '56.

(MERA 9:11)

1. Leningradskiy gosudarstvennyy universitet.

(Tantalum) (Selenium)





79-2-3/58

AUTHORS: Ariya, S. M.; Morozova, M. P.; Khuan Tszi-Tao; Vol'f, E.

TITE: The Enthalpy of Formation of Lithium, Magnesium and Zinc Arsenides

(Ental piya obrasovaniya arsenidov litiya, magniya i tsinka)

PERIODICAL: Zhurnal Obshchey Khimii, 1957, Vol. 27, No. 2, pp. 293-295 (U.S.S.R.)

ABSTRACT: The formation enthalpies of lithium, magnesium and zinc arsenides were experimentally established at -81.3 ± 2 , -96 ± 3 and -30.5 ± 3 kcal/g respectively. Numerous facts are cited indicating that the formation enthalpy value of arsine is in agreement with the data on the thermal stability of arsenides. Light appears to be a somewhat more exothermal compound than Light which is in conformity with the

fact of displacing the Sb by As from the combination with Li.

Card 1/2 There are 7 references, of which h are Slavic.

CIA-RDP86-00513R00010201 "APPROVED FOR RELEASE: Thursday, July 27, 2000

B-8 Thermodynamics, Thermochemistry, Equilibria, Physical-Chemical Analysis, Phase Transitions.

: Referat Zhuz Khimiya, No 1, 1958, 352 Abs Jour

C.M. Ariya, M.P. Morozova, S.A. Shchukarev. Author

: Enthalpies of Formation of Binary Compounds of Elements Inst Title

of Main Subgroup of V Group. Phenomenon of Secondary

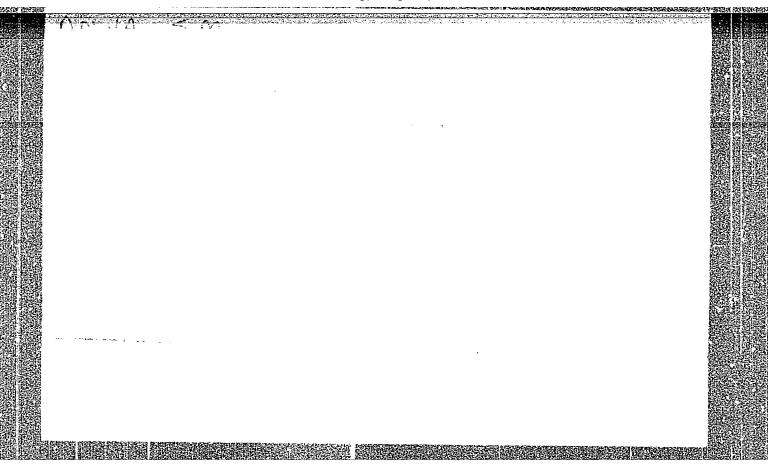
Periodicity.

Zh. obshch. khimii, 1957, 27, No 5, 1131-1136 Orig Pub

The phenomenon of the secondary periodicity (Biron Ye.V., Abstract

Zh. Russk. khim. o-va, 1915, 47, 964) of properties of elements of the main subgroups of the periodical system was investigated on the example of the course of enthalpy formations Δ H of elements of the main subgroup of the V group. The course of the A H changes is periodical in some cases and monotonous in other. In connection with the above, the valency states of atoms and the

Card 1/2



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ARIYA, S.M.; YEROFEYEVA, M.S.; MOCHALOV, G.P.

Magnetic susceptibility of strontium subnitride. Zhur.ob.khim.
27 no.7:1740-1743 J1 '57. (MIRA 10:10)

1. Leningradskiy gosudarstvennyy universitet.

(Strontium nitride--Magnetic properties)

HRIYIZ, S. M.

USSE/Physical Chemistry - Thermodynamics, Thermochemistry, Equilibria, Physical-Chemical Analysis, Phase Transitions.

Abs Jour: Referat. Zhurnal Khimiya, No 3, 1958, 7123.

Author : S.M. Ariya, Kan Kho-yn, Yu. Barbanel', G.M. Loginov.

Inst Title

: Enthalpy of Strontium Arsenide Sr3As2 Formation.

Orig Pub: Zh. obshch. khimii, 1957, 27, No 7, 1743-1745.

Abstract: Sr₃As₂ (I) was prepared by the interaction of the components at 400° in an evacuated glass tube and it was homogenized at 900° later. The pressure of As vapors on preparations of various composition was investigated by Knudsen's effusion method. The vapor pressure is minimum on I and it rises with the rise of As percentage in the preparations. The I formation enthalpy was determined from the data on I solubility in hydrochloric acid and on enthalpy magnitude of I interaction with

Card : 1/2

-11-

AUTHORS:

Ariya, S. M., Morozova, M. P.

SOV/79-28-10-1/60

TITLE:

Properties of Salt-Like Compounds of Variable Composition, and Ideas of Their Chemical Structures (Svoystva soleobraznykh soyedineniy peremennogo sostava i predstavleniya

o ikh khimioheskom stroyenii)

PERIODICAL:

Zhurnal obshchey khimii, 1958, Vol 28, Nr 10,

pp 2617 - 2623 (USSR)

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ABSTRACT:

One of the most important problems in general chemistry is whether a chemical compound has a constant or variable composition; which compounds represent considerable deviations from the stoichiometric composition, and how far these deviations are possible. The publications by N.S.Kurnakov exerted a considerable influence on the modern concept of the chemical structure. The rapid development of the x-ray structure analysis of solids made it possible to approach the rules proposed by Kurnakov. It was pointed out that the compounds of

variable composition radiographically represent subtraction, affiliation or substitution lattices (Refs 1-8). Typical

Card 1/3

affiliation or substitution lattices (Refs 1-8). Typical compounds of this type are FeO_{1+x} , TiO_{1+x} and VO_{1+x} which

Properties of Salt-Like Compounds of Variable Composition, and Ideas of Their Chemical Structures

SOV/79-28-10-1/60

are looked upon as defect structures of the NaCl type. The task of the present paper was to determine the dependence of some properties, first of all of the thermodynamic characteristic features, of the composition in some binary systems with compounds of variable composition. The investigation of the different types of dependences of some properties on the composition in these systems, as well as the thermodynamic investigation of the problem of the chemical structure of compounds of variable composition, lead to the conclusion that in the lattices of these compounds a segregation (of different power) of the atoms of the element of the substituting subgroup takes place with the atoms being in different states of valence. In short: The lattices of these variable compounds can have a submicro-unequal structure in different cases, i.e. remain monophase in the thermodynamical sense of the word. The part of the elements in this structure must depend on the nature of the compound on the temperature, as well as on the degree of deviation of these compounds

Card 2/3

Properties of Salt-Like Compounds of Variable Composition, and Ideas of Their Chemical Structures

SOV/79-28-10-1/60

with respect to the stoichiometric composition. There are 5 figures and 20 references, 13 of which are Soviet.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet (Leningrad

State University)

SUBMITTED:

December 31, 1957

Card 3/3

66247 sov/181-1-7-3/21 Ariya, S. M., Bogdanova, N. I. 24(3) 24.7700 Electrical Conductivity of Some Titanium and Vanadium Oxides AUTHORS: Fizika tverdogo tela, 1959, Vol 1, Nr 7, pp 1022-1026 (USSR) TITLE: Some time ago, it was supposed that some inconstant com-PERIODICAL: pounds, especially a series of oxides possess a submicroscopio inhomogeneous structure, e.g. that atoms of the trivalent ABSTRACT: titanium form accumulations, so-called "islets" in the lattice of titanium protoxide (TiO1+x). It was the intention of the author to explain whether the dependence of the electrical conductivity with regard to the composition in the system TiO1.00-TiO1.50 matches the above assumption. Conductivity measurements in vanadium oxides within the limits of VO_{1.50}-VO_{2.00} were performed, because within this zone numerous discrete intermediate compounds occur. The conductivity was investigated till 600°C by means of a potentiometer sonde. The results are reproducible in a satisfactory way taking account of the poly-crystalline sintered nature of the samples. The dependence of conductivity in vanadium oxides Card 1/3

Electrical Conductivity of Some Titanium and Vanadium Oxides

66247 SOV/181-1-7-3/21

on their composition (Fig 1) shows, that the conductivity of discrete intermediate compounds, in which the vanadium occurs in 3- and 4-valent formation, is different from the conductivity of a V₂O₃ and VO₂ mixture. The electrical conductivity of titanium oxide within the limits of TiO_{1.00}-TiO_{1.50}

equals almost the conductivity of a TiO_{1.00} and TiO_{1.50} mixture (Fig 2). The shape of the curve of the dependence of conductivity on the composition of that part within the homogeneous range of titanium protoxide (TiO_{1.00}-TiO_{1.20}), that is rich in oxygen, is very similar to the shape of the curve within the range of VO_{1.00}-VO_{1.67}. The latter corresponds to the heterogeneous range of the system V-O according to the radiographical data by Anderson and according to results of thermodynamical and magnetic investigations made by the authors(the thermodynamical investigations were shared by Yu. T. Popov). Consequently, the results of conductivity measurements are not contrary to the assumption of a submicroscopical inhomogeneous structure of the lattice

Card 2/3

√ 5 (2)

AUTHORS:

Vol'f, E., Ariya, S. M.

807/79-29-8-3/81

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TITLE:

Enthalpy of Formation of Vanadium Oxides

PERIODICAL:

Zhurnal obshchey khimii, 1959, Vol 29, Nr 8, pp 2470 - 2473

ABSTRACT:

The vanadium oxides necessary for the investigation were prepared in a special furnace at 1,600° by fusing the mixtures of the pulverized, hydrogenized vanadium and vanadium oxide which had been pressed into tablets. The analysis of the preparations obtained was made by determination of the weight gained in their oxidation to V205. The radiographic investigation of the pre-

parations showed that under these conditions the range of homogeneity of the vanadium suboxide lies within the interval VO 0,86 - VO 1,27. The heats of combustion of vanadium oxide were determined calorimetrically using small quantities (0,1-0,2 g) and an oxygen pressure of 42 atm. The gross composition of the combustion product was determined by the weight gained in oxidation. Results obtained in the determination of the heats of

Card 1/2

combustion (Q,p) of vanadium oxide and of metallic vanadium (with corrections) and the heats of formation of the various

Enthalpy of Formation of Vanadium Oxides

807/79-29-8-3/81

compositions computed from them are shown in table 1. The heats of formation hitherto determined of the vanadium oxides $VO_{\frac{1}{2}}, V_{2}O_{3}, VO_{2}$, and $V_{2}O_{5}$ differ considerably from each other. In table 2 the data obtained by the authors are compared with those given in publications. They correspond well to those by H. Siemonsen and to those suggested by the American Bureau of Standards if the sources of errors are taken into consideration. Thus, it was ascertained that the heat of formation of vanadium suboxide changes steadily with the composition, as is the case with titanium suboxide. This regularity is not in contradiction to the concept of the inhomogeneous submicroscopic structure of the lattice of some oxides of variable composition. There are 1 figure, 2 tables, and 9 references, 7 of which are Soviet.

ASSOCIATION:

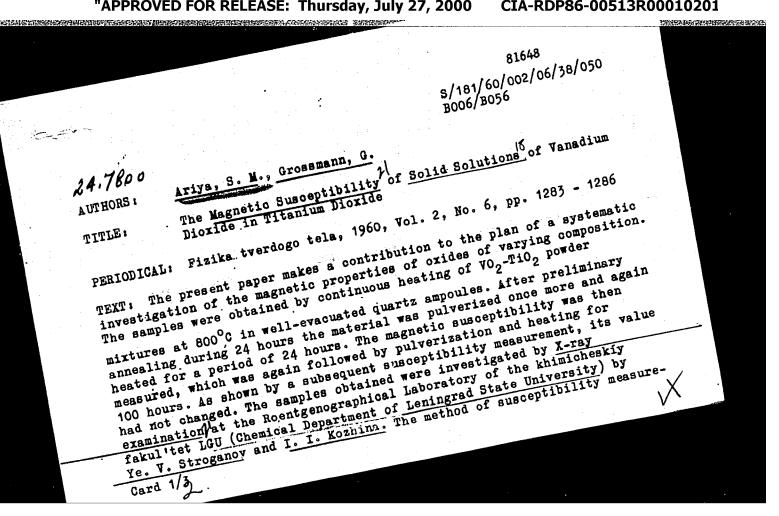
Leningradskiy gosudarstvennyy universitet (Leningrad State

University)

SUBMITTED:

July 11, 1958

Card 2/2



The Magnetic Susceptibility of Solid Solutions of Vanadium Dioxide in Titanium Dioxide

81648 S/181/60/002/06/38/050 B006/B056

ment is described in Ref. 1. Results are given in tables and diagrams. Fig. 1 shows the temperature dependence of χ within the range of 20-90°C for samples with different (30-100%) VO₂ content; the corresponding numerical values are given in Table 1. Table 2 gives the values of the susceptibilities (χ), of the constants of the Curie-Weiss law, and the effective magnetic moments for pure VO₂, pure TiO₂, and samples with 20 and 10 mole% VO₂. Fig. 2 shows the dependence of the paramagnetic component of χ per gram-atom of vanadium on the VO₂ concentration at different temperatures. The curves show that the suceptibility of solid VO₂-TiO₂ solutions is not additively composed of the susceptibilities of the components. Fig. 3 shows the temperature dependence of the reciprocal susceptibility of 4 samples of different composition. The results are briefly discussed, and are compared with those obtained by Rüdorff (Refs. 6,7). There are 3 figures, 2 tables, and 7 references: 1 Soviet, 4 German, 1 French, and 1 American.

Card 2/3

APPROVED FOR RELEASE: Thursday, July 27, 2000 CIA-RDP86-00513R000102010

84597

24,7900 (1055,1144,1160)

\$/181/60/002/010/020/051 B019/B056

AUTHORS:

Grossmann, G. and Ariya, S. M.

TITLE:

The Magnetic Susceptibility of a Solid Solution of Ferrous Oxide in Magnesium Oxide

PERIODICAL:

Fizika tverdogo tela, 1960, Vol. 2, No. 10, pp. 2477-2479

TEXT: The authors investigated the susceptibility of three preparations (FeO_{1.048}.0.487MgO, FeO_{1.080}.0.487MgO and FeO_{1.5}.0.487MgO) within the temperature range of from 700 to 900°C. The production of the preparations has been described earlier (Ref. 4). The magnetic susceptibility of ferrous oxide is a linear function of composition and it may be seen from the experimental results shown by the Fig. that the straight line, which characterizes the susceptibility of the two compositions FeO_{1+x} . MgO, on extrapolation up to the composition $\text{FeO}_{1.00}^{\circ}\text{O.487MgO}$ leads exactly to that value, to which the analogous straight lines for FeO_{1+x} and

Card 1/2

KOLBINA, Ye.M. [deceased]; BARRANEL', Yu.A.; NAZAROVA, M.V.; ARIYA, S.M.

Thermodynamics of lower cobalt sulfides. Vest. IGU 15 no.4:122-129
'60. (MRA 13:2)

(Gobalt sulfide) (Thermodynamics)

5,4100

77340

SOV/79-30-1-1/78

AUTHORS:

Bogdanova, N. I., Ariya, S. M.

TITLE:

Composition of Higher Oxides of Vanadium According to

Their Electric Conductivity Data

PERIODICAL:

Zhurnal obshchey khimii, 1960, Vol 30, Nr 1, pp 3-7

(USSR)

ABSTRACT:

The binary compounds whose structures apparently contain atoms of a metal in different valence states and whose compositions, consequently, cannot be expressed by using whole valence numbers, are termed "mixed" compounds. Vanadium oxides have been known to be among them since G. Anderson's X-ray investigations (Acta Chem. Scan., 8, 1599, 1954). He and other investigators found discrete "mixed" compound of VO_n composition with fractional n values such as 1.67, 1.75, 1.80, 1.84, 1.86,

1.87, 2.17; α -phase with n = 1.71-2.0; α' -phase with n = 2-2.23; α -phase with n = 1.67-1.79; α' -phase with

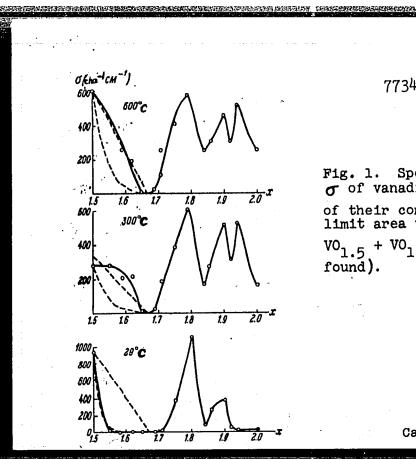
Card 1/6

Composition of Higher Oxides of Vanadium According to Their Electric Conductivity Data

77340 SOV/79-30-1-1/78

n up to 1.5. The authors examined the electric conductivity of the compounds, whose n varied from 1.5 to 2.0. The samples were prepared by a 4-stage treatment: (1) reduction of very pure V_2O_5 to V_2O_3 with thoroughly purified electrolytic H_2 at temperatures gradually raised up to 900° C; (2) annealing of $V_2O_3 + V_2O_5$ mixture under vacuum at 700° C for 10 hr, then at 900° C for 20 hr; (3) compression of the annealed and powdered mixture into tablets under $10,000 \text{ kg/cm}^2$ pressure; (4) sintering at $1,600^{\circ}$ C in a high-vacuum high-frequency furnace. The composition of the sintered tablets was determined according to the weight increase (oxygen consumption) on oxidation to V_2O_5 . The electric conductivities, measured with a potentiometer under vacuum at 20° C to 600° C, are shown in Fig. 1 and 2. The first 3 of the maxima and minima in Fig. 1 correspond to the known discrete compounds with n = 1.67, 1.80, 1.84,

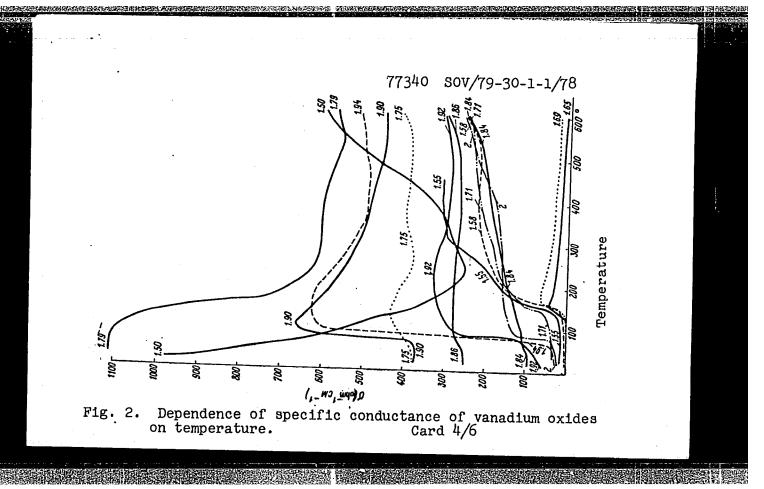
Card 2/6



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Fig. 1. Specific conductance σ of vanadium oxides as function of their composition (dotted lines limit area within which values of σ of $v_{1.5}$ + $v_{1.67}$ mixtures can be found).

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Composition of Higher Oxides of Vanadium According to Their Electric Conductivity Data

77340 SOV/79-30-1-1/78

while the origin of peaks at n = 1.90 and n = 1.94, and of dip at n = 1.92 is still unknown; perhaps they also indicate discrete compounds, the possibility of whose existence was mentioned by Anderson. The n values from 1.50 to 1.67 seem to correspond to 2 phase systems. No solid solution with n = 1.75 is indicated in the σ vs. composition curve, but absence of an abrupt change in the conductivity of VO_{1.75} at 160° C and essentially dif-

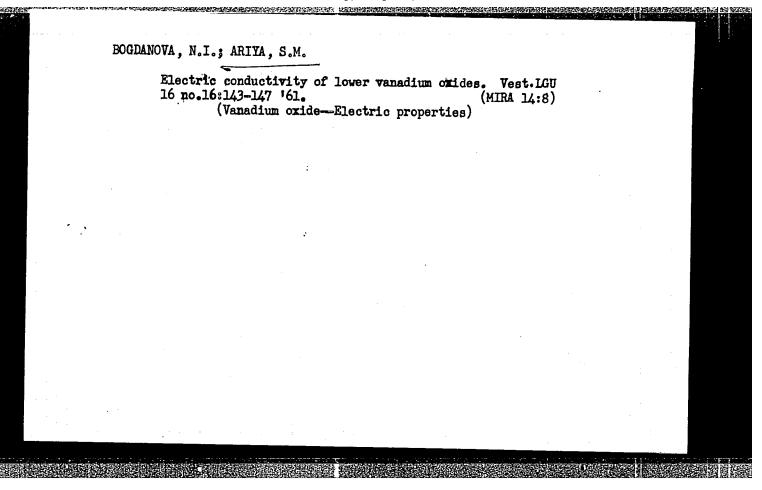
fering n=1.75 and n=1.79 curves in Fig. 2 point to existence of a discrete compound with n=1.75. Also, no compounds with n=1.86 and n=1.87 appear in the σ vs. composition curves. However, inspection of Fig. 2 permits one to assume a discrete compound or even 2 compounds at the interval from n=1.84 to n=1.88. There are 2 figures; 1 table; and 8 references, 3 German, 2 Soviet, 1 Danish, 1 Swiss, 1 Italian.

Card 5/6

ARIYA, S.M.; SOBOLEVA, M.S.

Lattice energies of oxides. Fiz.tver.tela 3 no.10:3157-3161 0
161. (MIRA 14:10)

1. Leningradskiy gosudarstvennyy universitet imeni A.A.Zhdanova.
(Oxides) (Crystal lattices)



21999 s/076/61/035/004/004/016 B106/B201

15.2142

Yerofeyeva, M.S., Lukinykh, N.L., and Ariya, S.M.

AUTHORS 3

Card 1/

Heat content of some titanium oxides at high temperatures

PERIODICAL: Zhurnal fizicheskoy khimii, v. 35, no. 4, 1961, 772 - 775 Several properties of compounds of a composition in the oxygen-rich part of the homogeneous titanium oxide phase (TiO1.00 - TiO1.20) appear to be evidence of the fact that these compounds have the lattice of TiO1.00 which contains submicroscopic inclusions of the composition TiO 1.50 (Ti₂0₃), statistically distributed at random. Because of the very small sizes of these inclusions, the whole system behaves as a homogeneous phase. If this assumption is correct, the heat capacity of such compounds must be equal to the heat capacity of a mixture of TiO and Ti203 of equal gross composition. In this connection, the authors examined the mean heat capacity of titanium oxides as a function of the composition at high temperatures. The titanium exides were prepared by annealing mixtures of titanium

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Heat content of some titanium ...

hydride and titanium dioxide in high vacuum at 1300°C. The composition of the oxides was determined from the weight increase in the oxidation to titanium dioxide in an aqueous oxygen flow at 1000 - 1100°C. The heat contents were determined by an apparatus resembling the one described in Ref. 5 (J.C. Southard, J. Amer. Chem. Soc., 62, 3112, 1941). For a test of the apparatus, the heat content of α -Al₂0₃ was measured at 200-800°C; the results were found to be in agreement with data available in the literature. The heat content of the titanium oxides was measured at 2200, 412°, 604°, and 809° C. The mean heat capacity of homogeneous compounds having a composition between TiO1.00 and TiO1.20 was found practically to coincide with the mean heat capacity of the mixture of TiO1.00 and TiO1.50 of equal gross composition. This result is not, however, explained by the fact that titanium ions are found side by side in the same form as in pure TiO1.00 and TiO1.50 in the lattice of the compounds concerned. In fact, experiments have revealed that the mean heat capacity of TiO1.67 (Ti305), in the lattice of which trivalent and tetravalent titanium ions are mani-Card 2/6

21999 S/076/61/035/004/004/018 B106/B201

Heat content of some titanium ...

festly present side by side, does not coincide with the mean heat capacity of a mixture of Ti,0, and TiO, of equal gross composition. Similarly, neither the mean heat capacity of Fe 304 coincides with the mean heat capacity of a mixture of FeO and Fe₂O₃ of equal gross composition. The reason for this is the structure sensitivity of the heat content. Ti(III) and Ti(IV) ions are in the lattice of TiO1.67 subjected to structural conditions other than in the lattices of TiO1.5 and TiO2, respectively. lar considerations apply to Ti(II) and Ti(III) ions in the lattice of compounds of a composition between TiO1.00 and TiO1.20. The coincidence of the mean heat capacity of these compounds with the mean heat capacity of a mixture of TiO and $\mathrm{Ti}_2\mathrm{O}_3$ may be explained by the fact that the atoms of trivalent titanium are concentrated in the lattice of $\text{TiO}_{1 \bullet 00}$ in the form of submicroscopic inclusions. The Ti(III) atoms and also the oxygen atoms bound with them would have the same environment as in the lattice of Ti,03, and would therefore contribute to the heat capacity of TiO1+x an Card 3/6

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Heat content of some titanium

amount as much as corresponds to the heat capacity of the same amount of Ti203. The number of Ti(III) atoms concentrated in the submicroscopic inclusions is as yet still unknown; this problem will be dealt with in a following paper. The fact is stressed here that while the abovementioned assumption explains satisfactorily the additive composition of the heat capacity of compounds between TiO1.00 and TiO1.20 by the heat capacities of TiO1.00 and TiO1.50, it cannot be taken as a proof that compounds of the structure TiO are actually submicroscopically heterogeneous. has been found that the form of the dependence of the mean heat capacity of TiO1+x compounds on the composition changes in the point of the stoichiometric composition (TiO1,00). Similar changes have been observed also by other authors in the dependence of the formation enthalpies and of the volumes of the formulas expressed in g on compounds of the type Tio, There are 3 figures and 7 references: 6 Soviet-bloc and 1 non-Soviet-bloc. The reference to the English language publication reads as Card 4/6

21999 \$/076/61/035/004/004/018 B106/B201

Heat content of some titanium ...

follows: J.C. Southard, J. Amer. Chem. Soc., 62, 3112, 1941.

ASSOCIATION:

Leningradskiy gosudarstvennyy universitet im. A.A. Zhdanova (Leningrad State University imeni A.A.

Zhdanov)

SUBMITTED:

July 13, 1959

Card 5/6

S/181/62/004/010/044/063 B102/B112

AUTHORS:

Ariya, S. M., and Golomol'zina, M. V.

TITLE:

Infrared spectra of titanium and vanadium oxides in the

crystalline state

PERIODICAL:

Fizika tverdogo tela, v. 4, no. 10, 1962, 2921-2924

TEXT: Since it is not possible to determine roentgenographically whether or not the vacancies and complexes of vacancies in lower crystalline titanium and vanadium oxides are randomly distributed, the effect of the oxygen content of these lower oxides on the IR spectra is studied. The spectra were taken with an NKC-14 (IKS-14) double-ray spectrometer. The oxides were added to pure KBr in a concentration of 0.1% and the mixture was pressed to tablets. The compounds TiO_{0.94}, TiO_{1.00}, TiO_{1.17}, TiO_{1.5}, VO_{0.95}, VO_{1.15}, VO_{1.28} and VO_{1.5} were studied. For titanium oxides, the most important lines and the general form of the IR absorption spectrum were found to agree with those of Ti₂O₃, except that in the lower oxides the lines at 490 and 520 cm⁻¹ are shifted towards 470 and Card 1/2

Infrared spectra of titanium

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514 cm⁻¹ respectively. The distinct 1080 cm⁻¹ line is common to all oxides. For vanadium the lines of the lower oxides are consistent with those of V₂O₃. These spectra, X-ray pictures, and a comparison with the IR spectra of other metal oxides indicate that all oxides are TiO_{1±x} and VO_{1±x} mixtures of pure metal and Me₂O₃. Blokhin and Shuvayev (Izv. AN SSSR, ser. fiz.26,1962), also, found that the TiO spectrum can be regarded as a superposition of the Ti and Ti₂O₃ spectra, but they gave a different explanation. There are 5 figures.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet (Leningrad State University)

SUBMITTED:

June 12, 1962

Card 2/2

AUTHORS: Morozova, M. P., Khripun M. K. and Ariya, S. M. S/079/62/032/007/001/007 TITLE: 1032/1232

The enthalpy of carbides and oxycarbides of titanium PERIODICAL:

Zhurnal obshchei khimii, v. 32, no. 7, 1962, 2072-2076 TEXT: The enthalpy of three titanium carbides, ranging in composition from TiC_{0.79}. to TiC_{1.00}, and of six different titanium oxycarbides ranging between the compositions TiC_{0.15}O_{0.096}, TiC_{0.42}O_{0.118} and TiC_{0.74} O_{0.059}, was calculated from the heat of combustion of these compounds, determined calorimetrically, and from the known enthalpies of TiO₂ and CO₂. The value of the enthalpy of formation of TiC is given as -55 ± 0.3 Kcalories per mole. This result is compared with that given by Humphry. The dependence of the enthalpy on the index at the C atom in the composition interval TiC_{1.00}-TiC_{0.79} is found to be linear. The

enthalpy of oxycarbides TiC_xO_y is found to be equal to the sum of the enthalpies of TiC_x and TiO_y. Hence it is inferred that the coexistence of Ti-C and Ti-O bonds in the lattice of oxycarbides has practically no effect on the energy of these bonds. There is 1 figure and 3 tables. English-language references read: K. K. Kelley, U. S. Bur. Mines Rept. Invest, No. 5316, 33 (1957). J. Humphry, J. Am. Chem. Soc., 73, 2261 (1951). ASSOCIATION: Leningradskii gosudarstvennyy universitat (Leningrad State University)

SUBMITTED:

Card 1/1 ,

ARIYA, S.M., POPOV, Yu., Structure of titanium and vanadium monoxide lattices. Zhur.ob. khim. 32 no.7:207;~2081 Jl. *62. (MIRA 15:7)

1. Leningradskiy gosudarstvennyy universitet, (Titanium exides) (Vanadium exides) (Crystal lattices)

ARIYA, S.M.; MOROZOVA, M.P.

Equilibrium of vanadium oxides with CO2/CO mixtures. Zhur.ob.khim.
32 no.7:2081-2083 Jl *62. (MIRA 15:7)

1. Leningradskiy gosudarstvennyy universitet.
(Vanadium oxides) (Carbon oxides)

ARITA, S.M.; GOLOMOLZINA, M.V.

Infrared spectra of titanium and vanadium oxides in the crystalline state. Fiz.tver.tela 4 no.10:2921-2924 0 '62.

1. Leningradskiy gosudarstvennyy universitet.
(Titanium oxide crystals-Spectra)
(Vanadium oxide crystals-Spectra)

ARIYA, S.M.; BRACH, B.Ya.

Electroconductivity of iron oxide at high temperatures. Fiz. tver tela 5 no.12:3496-3499 D '63. (MIRA 17:2)

1. Leningradskiy gosudarstvennyy universitet.

